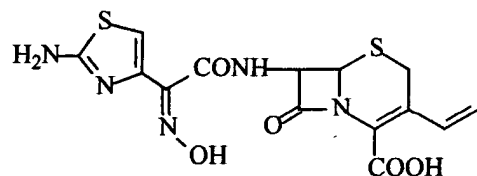




**WE CLAIM:**

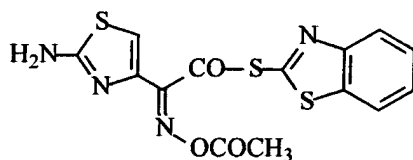
1. (currently amended): A process for preparing a cephalosporin cefdinir of Formula II



Formula II

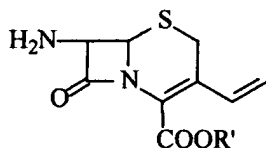
comprising the steps of:

reacting O-acetyl thioester of Formula I



Formula I

with a compound of Formula III in the presence of a base in suitable solvent

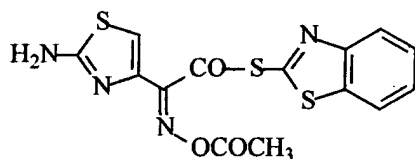


Formula III

wherein R' [[= H,]] represents H or any carboxyl protecting group or silyl group,  
converting to cefdinir by the removal of protecting group or groups.

2. (original): The process according to claim 1 wherein the said base can be organic base or an inorganic base.
3. (currently amended): The process according to claim 1 wherein the said organic base is an amine selected from the group consisting of triethylamine, N,N-diisopropylethylamine, ~~tributylamine~~ tri n-butylamine.
4. (original): The process according to claim 1 wherein the said inorganic base is selected from the group consisting of sodium carbonate, sodium bicarbonate and mixtures thereof.
5. (original): The process according to claim 1 wherein the said solvent is selected from the group consisting of water, tetrahydrofuran, methylene dichloride and mixtures thereof.
6. (currently amended) The process according to claim 1 wherein the said reacting step is conducted at a temperature between 10°C and 25°C.
7. (currently amended) The process according to claim 1 wherein the said carboxyl protecting group is selected from the group consisting of p-methoxybenzyl, p-nitrobenzyl, ~~and diphenylmethyl~~ and trimethyl silyl .

8. (currently amended): The process according to claim 1 wherein the said to prepare  
O-acetyl thioester of Formula I



Formula I

- is prepared by a process which comprises of  
condensing (Z)-2-(2-amino-4-thiazolyl)-2-acetyloxyiminoacetic acid with  
bis(benzothiazol-2-yl)disulphide in the presence of triphenylphosphine and a  
base in a suitable solvent.
9. (original): The process according to claim 8 wherein the said base is selected from  
the group consisting of tributylamine, triethylamine and mixtures thereof.
10. (original): The process according to claim 8 wherein the said solvent is selected  
from the group consisting of methylene chloride, chloroform, tetrahydrofuran,  
acetonitrile and mixtures thereof.
11. (original): The process according to claim 8 wherein the said reacting step is  
conducted at a temperature between 0°C and 35°C.